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HOUSE BILL 1202

48TH LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 2007

INTRODUCED BY

Irvin Harrison

AN ACT

RELATING TO PUBLIC HEALTH; PROVIDING FOR TREATMENT OF CERTAIN
DISEASES WITHOUT A PRACTITIONER-PATIENT RELATIONSHIP OR AN
INTERVENING MEDICAL EVALUATION.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:

Section 1. Section 26-1-16 NMSA 1978 (being Laws 1967,
Chapter 23, Section 16, as amended) is amended to read:

"26-1-16. DANGEROUS DRUGS--CONDITIONS FOR SALE--
PRESCRIPTION REFILLING--LIMITATIONS.--

A. It is unlawful for [~~any~~] a person to sell,
dispose of or possess any dangerous drugs, except:

(1) manufacturers, wholesalers or
distributors, their agents or employees licensed by the board
to ship dangerous drugs into the state; or

(2) distributors, wholesalers, hospitals,

underscored material = new
[bracketed material] = delete

1 nursing homes, clinics or pharmacies and other authorized
2 retailers of dangerous drugs in this state licensed by the
3 board, and appropriate records of dangerous drugs receipt and
4 disposition are kept. These records shall be open to
5 inspection by any enforcement officer of this state.

6 B. Practitioners licensed in this state may
7 prescribe, provide samples of and dispense any dangerous drug
8 to a patient where there is a valid practitioner-patient
9 relationship. A record of all such dispensing shall be kept
10 showing the date the drug was dispensed and bearing the name
11 and address of the patient to whom dispensed. It is the duty
12 of every licensed physician, dentist, veterinarian, pharmacist
13 or person holding a limited license issued under Subsection B
14 of Section 61-11-14 NMSA 1978, when dispensing any dangerous
15 drug, to mark on the dispensing container the name of the
16 patient, the date dispensed, the name and address of the person
17 dispensing the drug, the name and strength of the drug,
18 expiration date where applicable, adequate directions for use
19 and the prescription number when applicable. All official
20 compendium requirements for the preservation, packaging,
21 labeling and storage of dangerous drugs are applicable where
22 drugs are held for dispensing to the public, whether by a
23 pharmacy, clinic, hospital or practitioner.

24 C. Notwithstanding the provisions of Subsection B
25 of this section, a practitioner licensed in this state may

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1 prescribe, provide samples or dispense a dangerous drug to a
2 sex partner of a person with a sexually transmitted disease
3 without a valid practitioner-patient relationship or an
4 intervening medical evaluation. The New Mexico medical board
5 shall promulgate rules to carry out the provisions of this
6 subsection, including identification of the diseases that may
7 be treated pursuant to this subsection.

8 ~~[G.]~~ D. Pharmacists are prohibited from selling or
9 disposing of ~~[any]~~ a dangerous drug except on prescription of a
10 practitioner and except as such sale or possession is
11 authorized under Subsection A of this section. It is the duty
12 of all pharmacists to keep an accurate record of all disposals,
13 which record shall be open to inspection by ~~[any]~~ an
14 enforcement officer of this state.

15 ~~[D.]~~ E. No enforcement officer having knowledge by
16 virtue of ~~[his]~~ office of ~~[any]~~ a prescription, order or record
17 shall divulge such knowledge except in connection with a
18 prosecution or proceeding in court or before a licensing or
19 registration board or officer, to which prosecution or
20 proceeding the person to whom such prescriptions, orders or
21 records relate is a party.

22 ~~[E.]~~ F. It is unlawful, except as otherwise
23 authorized under Subsection A of this section or the Controlled
24 Substances Act and except for the college of pharmacy of the
25 university of New Mexico or a public health laboratory, for

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1 ~~[any]~~ a person to possess any dangerous drug unless such
2 substance has been dispensed to ~~[him]~~ the person either
3 directly by a practitioner or on a prescription.

4 ~~[F-]~~ G. All records required to be kept under the
5 provisions of the New Mexico Drug, Device and Cosmetic Act
6 shall be preserved for a period of three years, provided that
7 records requirements do not apply to the administration of a
8 drug to a patient upon whom the practitioner personally
9 attends, and provided that records of controlled substances
10 shall be kept in accordance with the provisions of the
11 Controlled Substances Act.

12 ~~[G-]~~ H. No prescription may be lawfully refilled:

13 (1) if it is marked by the issuing
14 practitioner as not to be refilled;

15 (2) when the practitioner indicates a specific
16 number of refills or a specific period of time, on the original
17 prescription calling for a dangerous drug, it may be refilled
18 the number of times or for the period of time indicated;
19 provided the date of refill, the initials of the pharmacist
20 refilling the prescription and the amount of drug dispensed, if
21 it differs from the amount called for on the original
22 prescription, is recorded on the original prescription and
23 provided a prescription issued for drugs controlled by the
24 Controlled Substances Act shall comply with that act;

25 (3) when the practitioner does not indicate

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1 refill instructions on the original prescription calling for a
2 dangerous drug, unless:

3 (a) the practitioner is contacted
4 orally, by telephone, telegraph or other means of communication
5 for instruction; and

6 (b) if authorization to refill is given
7 the pharmacist, the following information will be immediately
8 transferred to the original prescription: 1) date; 2) name of
9 person authorizing the refill; 3) pharmacist's initials; and 4)
10 amount dispensed if different [~~than~~] from the amount indicated
11 on the original prescription;

12 (4) when the practitioner indicates on the
13 original prescription calling for dangerous drugs that it may
14 be refilled "prn", the pharmacist may refill it within the
15 limits of the dosage directions for a period of twelve months;
16 provided the date of refilling and the initials of the
17 pharmacist are recorded on the original prescription. At the
18 expiration of the twelve-month period, the practitioner must be
19 contacted for a new prescription; provided that this is not to
20 be construed to apply to those drugs regulated by the
21 Controlled Substances Act; and

22 (5) the board may adopt and promulgate
23 regulations to permit the use of computer systems for the
24 storage and retrieval of prescriptions, records for the purpose
25 of refilling prescriptions, receipt records, drug distribution

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1 records, drug withdrawals from stock, drug compounding records,
2 drug disposition records and drug disposal records.

3 ~~[H.]~~ I. Nothing in this section shall prevent the
4 owner of livestock or ~~[his]~~ the owner's consignee or ~~[their]~~
5 employees ~~[to be]~~ from being in possession of drugs for ~~[their]~~
6 use in performing routine, accepted livestock management
7 practices in the care of livestock belonging to the owner, and
8 if the drugs are labeled as being restricted to animal use
9 only; provided that if such drugs bear the legend: "CAUTION:
10 federal law restricts this drug to use by or on the order of a
11 licensed veterinarian", the drugs may be used or distributed
12 only as provided in Subsection A of Section 26-1-15 NMSA 1978."